## **CLAIMS**

- 1. Pharmaceutical compositions having a stimulating effect on the proliferation of NK cells, comprising an effective amount of at least an antibody selected in the group comprising an anti-NCR antibody such as anti-NKp30 antibody or anti-NKp46 antibody, or both, or an immuno-reactive fragment thereof, and a cytokine selected in the group comprising interleukins such as IL2, IL12, IL15, IL21 or a combination thereof, in association with a pharmaceutically acceptable carrier, said antibody(ies) and cytokine(s) being administered together or separately to a subject.
- 2. The pharmaceutical compositions of claim 1, wherein said anti-NKp30 antibody and/or anti-NKp46 antibody are used in admixture with IL2.
- 3. The pharmaceutical compositions of claim 1, wherein said anti-NKp30 antibodies are isolated antibodies or antigen binding fragment thereof which specifically bind to a polypeptide selected from the group consisting of SEQ ID N°1, SEQ ID N°2, SEQ ID N°3, SEQ ID N°4, or an immunogenic fragment thereof, and SEQ ID N°5.

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- 4. The pharmaceutical compositions of claim 3, wherein said antibodies specifically bind to polypeptide having SEQ ID N°1.
- 5. The pharmaceutical compositions of claim 1, wherein said anti-NKp30 and/or anti-NKp46 antibodies are monoclonal antibodies, affinity, chimerized or humanized antibodies and more preferably humanized mouse monoclonal antibodies or of human origin.
- 6. The pharmaceutical compositions of claim 5, wherein said anti-NKp30 monoclonal antibody is produced by hybridoma strain I-2576.

- 7. The pharmaceutical compositions of claim 1, comprising antibody fragments, said fragments being essentially Fab, F(ab')<sub>2</sub>, and Fv fragments and CDR grafted humanized monoclonal antibodies.
- 8. The pharmaceutical compositions of claim 1, which are administered by various routes, including intradermal, intramuscular, intraperitoneal, intravenous, or subcutaneous injection, intranasal route and the chirurgical route.
- 9. The pharmaceutical compositions of claim 1, which are under the form of tablet, powder, pastes, patches, granules, microgranules, nanoparticules, colloid solution, aqueous solution, injectable solutions, sprays, liposomes.
- 10. The pharmaceutical compositions of claim 1, when used for daily subcutaneous injection, comprising from 1 ng to 100mg/kg (body weight) of antibodies, and lower than 1 million units/square meters/day of cytokine(s), for prevention, palliation, therapy e.g. of melanoma, Chronic Myeloid Leukemia, Acute Myeloid Leukemia, Lymphomas, Multiple Myeloma, hepatocarcinoma, lung adenocarcinoma, Neuroblastoma and for anti-microbial prevention, palliation and therapy.
  - 11. A method for stimulating the proliferation of NK cells which comprises contacting NK cells with an effective amount of a pharmaceutical composition according to claim 1.

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12. A method according to claim 11 comprising one or several injections of an effective amount of at least an antibody selected in the group comprising an anti-NCR antibody such as anti-NKp30 antibody or anti-NKp46 antibody, or both, or an immuno-reactive fragment thereof, and, repeated injections of a cytokine selected in the group comprising interleukins such as IL2, IL12, IL15,

- IL21 or a combination thereof, during 5-10 days, said cytokine(s) being first injected on the same day as the first injection of antibodies.
- 13. The method of claim 12, comprising one or two injections/day of cytokine(s)5 by subcutaneous route.
  - 14. The method of claim 11, wherein said interleukine is IL-2 and is injected subcutaneously at daily doses below 1 million units/m² for 5 to 10 days.
- 15. The use of the pharmaceutical compositions of claim 1 in the manufacture of a drug for prevention, palliation, therapy e.g., of melanoma, Chronic Myeloid Leukemia, Acute Myeloid Leukemia, Lymphomas, Multiple Myeloma, hepatocarcinoma, lung adenocarcinoma, Neuroblastoma and for anti-microbial prevention, palliation and therapy.